

CASE REPORT

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Successful use of recombinant activated factor VII (eptacog alfa, Novoseven®) in a refractory bleeding during pediatric cardiac surgery: a case report

Mariko Ishihara^{1*} , Yoshikazu Miyamoto², Akihiro Taniguchi² and Keiko Kinouchi²**Abstract**

A 2-year-and-7-month-old boy underwent an emergent reconstruction surgery of the right ventricle-pulmonary artery (RV-PA) conduit. Although he was successfully weaned from cardiopulmonary bypass (CPB) after reconstruction of the RV-PA conduit, the bleeding continued despite the massive transfusion of red blood cell (RBC), fresh frozen plasma (FFP), and platelets. Because of persistent bleeding and abnormal coagulation laboratory results, we administered the recombinant activated factor VII (rFVIIa), which was not approved for use in the treatment of post-CPB coagulopathy. After administration of rFVIIa, his coagulation data dramatically improved, the bleeding decreased, and the operation was able to be finished.

Keywords: Pediatric cardiac surgery; Refractory bleeding; rFVIIa; Acquired hemophilia A

Background

There have been increasing numbers of reports of “off-label” use of the recombinant activated factor VII (rFVIIa) to achieve hemostasis in various situations including traumatic intractable bleeding [1, 2], cardiac surgery [3, 4], and obstetrical massive bleeding [4]. However, there are few data regarding the appropriate use of rFVIIa, especially in pediatric patients. Here, we report a case of successful administration of rFVIIa for refractory post-cardiopulmonary bypass (CPB) bleeding in a child undergoing repeated cardiac surgery.

Case presentation

A 2-year-and-7-month-old boy with 22q11.2 deletion syndrome (height, 86 cm; body weight, 10 kg) was scheduled for an emergent surgery to reconstruct the right ventricle-pulmonary artery (RV-PA) conduit to remove its compression on his right coronary artery, causing the right ventricular failure. He was associated

with tetralogy of Fallot, pulmonary atresia, major aorto-pulmonary collateral arteries, and right aortic arch. He underwent a systemic-pulmonary shunt (left subclavian artery to central pulmonary artery) at 1 month of age and right unifocalization and right Blalock-Taussig shunt at 1 year and 3 months of age. When he was 2 years and 7 months old, he underwent RV-PA conduit and pulmonary artery plasty. After the procedure, because of the poor contraction of the right ventricle, weaning from CPB failed and extracorporeal membrane oxygenation (ECMO) was instituted, from which he was weaned on postoperative day 11.

Twenty-two days after surgery, cardiac catheterization revealed that his right coronary artery was compressed by the RV-PA conduit, and the emergent reconstitution of the RV-PA conduit was scheduled on the same day.

Anesthesia was induced with 20 µg of fentanyl and 10 mg of rocuronium, and repositioning of the RV-PA conduit was performed under CPB. He was successfully weaned from CPB with the support of inhaled nitric oxide (20 ppm), adrenaline (200 ng/kg/min), dopamine (10 µg/kg/min), and dobutamine (10 µg/kg/min). After reversal of heparin using protamine, persistent

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hemorrhage over 40 ml/kg/h continued for 3 h despite continuous massive administration of red blood cell (RBC), fresh frozen plasma (FFP), and platelets. At that time, abnormal coagulation tests were noted with prothrombin time (PT) of 36 % and international normalized ratio of PT (PT-INR) of 1.73, activated partial thromboplastin time (APTT) of 173 s, and fibrinogen was 172 mg/dl. The platelet count was 97,000/ μ L. As a rescue treatment to reduce bleeding and to correct his coagulation abnormality, a single dose of 100 μ g/kg of rFVIIa was intravenously administered. One hour later, coagulation tests were almost normalized as shown in Table 1 and the hemorrhage decreased to 5–10 ml/kg/h, and the operation was finished with the open sternum. He was transferred to the Pediatric Intensive Care Unit (PICU).

CPB time was 340 min, and his body temperature during CPB was 32 °C. Operation time was 639 min, and anesthesia time was 688 min. Total blood loss was 1480 ml, and required blood products were RBC 690, FFP 725, and platelets 250 ml, respectively.

Five hours after the administration of rFVIIa, coagulation data were comparable to the preoperative values (Table 1) and the total bleeding dose was less than 1 ml/kg/h. No additional rFVIIa was necessary. No thromboembolic events were noted after its administration.

Fifty-two days after the second surgery, his sternum and skin were able to be closed, and he was discharged from PICU on day 157.

Discussion

We reported the use of rFVIIa in a child complicated with persistent bleeding after cardiac surgery under CPB. Because massive transfusion of FFP and platelets failed to control post-CPB bleeding, we decided to use rFVIIa though we knew that post-CPB coagulopathy was not a licensed indication. In this case, rFVIIa was administrated without previous parental informed consent for its use. The administration of rFVIIa dramatically improved his coagulation data and reduced blood loss.

Recombinant FVIIa is a hemostatic agent developed for the treatment of hemophilia A or B with inhibitors

(antibodies) to coagulation factor VIII or IX. Currently, it is also approved to patients with congenital factor VII deficiency or the patients who have acquired antibodies to factor VIII, which is called as acquired hemophilia A. Furthermore, there have been increasing numbers of reports describing the off-label use of rFVIIa, which was effective in controlling refractory bleeding in nonhemophilic patients [1–4].

Recombinant FVIIa is considered to work by two mechanisms (Fig. 1): tissue factor (TF)-dependent and independent mechanisms. In the TF-dependent mechanism, FVIIa/TF complexes activate coagulation factors IX and X and activated factor X combines with factor V, resulting in an increase of thrombin generation. In the TF-independent mechanism, FVIIa can directly activate factor X on the platelet surface without the presence of TF. These mechanisms activate and aggregate platelets, resulting in formation of stabilized clots [5].

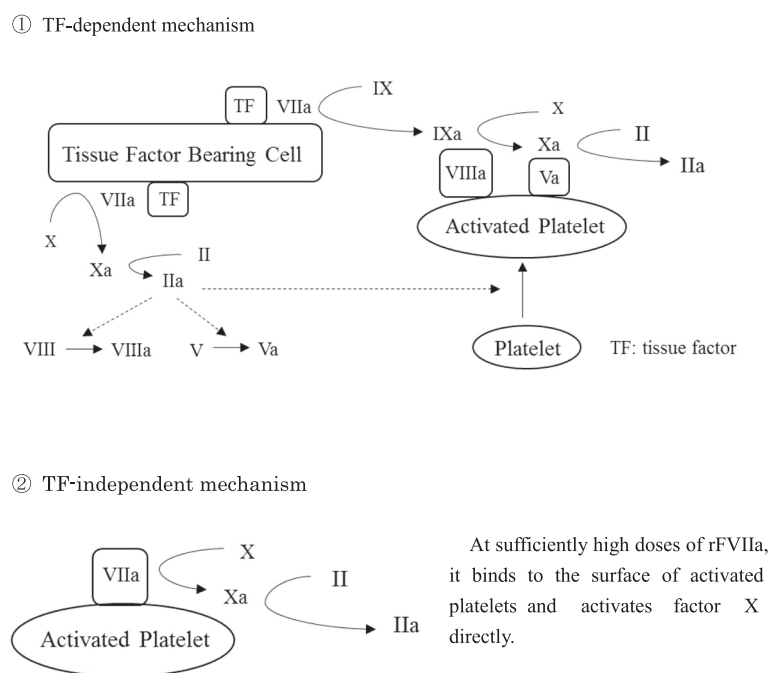
Uncontrolled bleeding after CPB is a critical problem especially in young children undergoing complex cardiac repair. The proposed mechanisms of the post-CPB coagulopathy in these young patients include (1) large circuit priming volume compared to the circulating blood volume, leading to dilution of platelets and coagulation factors; (2) long, complex cardiac repairs with extensive suture lines; (3) effects of hypothermia on both plasma coagulation factors and platelet function; (4) consumption of coagulation factors due to tissue injury, contact activation, and transfusion of suctioned pericardial blood; and (5) residual heparin or other anticoagulants [3, 5]. Adding to the repeated procedures, our patient was on the ECMO until 11 days before and underwent cardiac catheterization a few hours before the surgery in the unstable hemodynamic and debilitating state. These conditions might have further influenced the coagulation derangement, as partly shown by the abnormal data of PT, PT-INR, and APTT before surgery (Table 1).

There is no consensus on how to manage post-CPB coagulopathy in pediatric patients. The transfusion of platelets is the initial treatment for ongoing bleeding after adequate heparin neutralization in many

Table 1 Coagulation profile before and after administration of rFVIIa

		Normal value	Before surgery	Prior to rFVIIa	1 h after rFVIIa	5 h after rFVIIa
Platelet	/ μ L	130–400 $\times 10^3$	438,000	97,000	110,000	218,000
PT	%	80–120	53	36	>120	81
PT-INR		0.9–1.1	1.37	1.73	0.82	1.10
APTT	s	25–45	64	173	44	32
Fibrinogen	mg/dl	180–360	308	172	229	228

rFVIIa recombinant activated coagulation factor VII, PT prothrombin time, PT-INR international normalized ratio of PT, APTT activated partial thromboplastin time



Written informed consent was obtained from the parent of the patient for publication of this case report. A copy

of the written consent is available for review by the Editor of this journal.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

MI: data collection, literature search, manuscript drafting and anesthesiologist involved in the case. YM: manuscript drafting and anesthesiologist involved in the case. AT: manuscript drafting. KK: manuscript drafting and supervision. All authors read and approved the final manuscript.

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References

1. Yao D, Li Y, Wang J, Yu W, Li N, Li J. Effects of recombinant activated factor VIIa on abdominal trauma patients. *Blood Coagul Fibrinolysis*. 2014;25:33–8.
2. Brown CV, Sowery L, Curry E, Valadka AB, Glover CS, Grabarkewitz K, et al. Recombinant factor VIIa to correct coagulopathy in patients with traumatic brain injury presenting to outlying facilities before transfer to the regional trauma center. *Am Surg*. 2012;78:57–60.
3. Guzzetta NA, Russell IA, Williams GD. Review of the off-label use of recombinant activated factor VII in pediatric cardiac surgery patients. *Anesth Analg*. 2012;115:364–78.
4. Zatta A, Mcquiltan Z, Kandane-Rathnayake R, Isbister J, Dunkley S, Mcneil J, et al. The Australian and New Zealand Haemostasis Registry: ten years of data on off-licence use of recombinant activated factor VII. *Blood Transfus*. 2015;13:86–99.
5. Barua A, Rao VP, Ramesh B, Barua B, El-Shafei H. Salvage use of activated recombinant factor VII in the management of refractory bleeding following cardiac surgery. *J Blood Med*. 2011;2:131–4.
6. Hornykewycz S, Odegard KC, Castro RA, Zurakowski D, Pigula F, DiNardo JA. Hemostatic consequences of a non-fresh or reconstituted whole blood small volume cardiopulmonary bypass prime in neonates and infants. *Paediatr Anaesth*. 2009;19:854–61.
7. Pychyńska-Pokorska M, Pagowska-Klimek I, Krajewski W, Moll JJ. Use of recombinant activated factor VII for controlling refractory postoperative bleeding in children undergoing cardiac surgery with cardiopulmonary bypass. *J Cardiothorac Vasc Anesth*. 2011;25:987–94.
8. Levi M, Levy JH, Andersen HF, Truloff D. Safety of recombinant activated factor VII in randomized clinical trials. *N Engl J Med*. 2010;363:1791–800.
9. Dominguez TE, Brown KL. Activated recombinant factor VII and extracorporeal membrane oxygenation: everything is all right, nothing is wrong? *Pediatr Crit Care Med*. 2010;11:155–6.

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